

(PCT Article 36 and Rule 70)

REC'D 26 MAR 2004

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Applicant's or agent's file reference AXP/PG4787		FOR FURTHER ACTION	See Notification of Transmittal of International Preliminary Examination Report (Form PCT/IPEA/416)
International application No. PCT/EP 03/03347		International filing date (day/mor 27.03.2003	hth/year) Priority date (day/month/year) 28.03.2002
		both national classification and IPC	
C07D4	13/12		
Applicant			
Applicant GLAXO	: GROUP LIMITED et al.		
			·
1 76			
1. Th Au	is international preliminary exa thority and is transmitted to the	mination report has been prepa e applicant according to Article 3	red by this International Preliminary Examining 6.
i			
2. Thi	is REPORT consists of a total	of 7 sheets, including this cover	sheet.
			f the description, claims and/or drawings which have is containing rectifications made before this Authority
T .	(300 maio / 0.70 and occitor	n oor of the Administrative instru	ictions under the PCT).
The	ese annexes consist of a total of	of sheets.	
3. This	s report contains indications re	lating to the following items:	
1	☑ Basis of the opinion		
11	☐ Priority		
Ш	Non-establishment of o	Opinion with regard to novelty, in	ventive step and industrial applicability
IV	☑ Lack of unity of invention	on	remove step and industrial applicability
V	☑ Reasoned statement u		to novelty, inventive step or industrial applicability;
VI	☐ Certain documents cite		<i>,</i> •
IIV	Certain defects in the in	nternational application	
VIII		n the international application	
Date of sub	mission of the demand		
Date of Sup	mission of the demand	Date of c	ompletion of this report
30.09.2003		25.03.2	004
Name and malling address of the international preliminary examining authority:			d Officer
European Patent Office			South Marie Marie
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l. Ba	sis	of	the	re	oort
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1. With regard to the **elements** of the international application (Replacement sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this report as "originally filed" and are not annexed to this report since they do not contain amendments (Rules 70.16 and 70.17)):

	D	Description, Pages				
	1-	84	as originally filed			
	CI	aims, Numbers				
		22	as originally filed			
_			•			
2	. W lar	ith regard to the lang nguage in which the in	uage, all the elements marked above were available or furnished to this Authority in the nternational application was filed, unless otherwise indicated under this item.			
	Th	ese elements were a	vailable or furnished to this Authority in the following language: , which is:			
		the language of a to	anslation furnished for the purposes of the international search (under Rule 23.1(b)).			
		the language of pul	plication of the international application (under Rule 48.3(b)).			
		the language of a tr Rule 55.2 and/or 55	anslation furnished for the purposes of interpotional proliminary and in the purpose of the purpose o			
3.	. With regard to any nucleotide and/or amino acid sequence disclosed in the international application, the international preliminary examination was carried out on the basis of the sequence listing:					
			ernational application in written form.			
			ne international application in computer readable form.			
	furnished subsequently to this Authority in written form.					
		furnished subseque	ntly to this Authority in computer readable form.			
		The statement that the listing has been furn	he information recorded in computer readable form is identical to the written sequence ished.			
4.	The amendments have resulted in the cancellation of:					
		the description,	pages:			
		the claims,	Nos.:			
		the drawings,	sheets:			
5.	☐ This report has been established as if (some of) the amendments had not been made, since they have been considered to go beyond the disclosure as filed (Rule 70.2(c)).					
		(Any replacement sh report.)	eet containing such amendments must be referred to under item 1 and annexed to this			
6.	Add	itional observations, i	f necessary:			

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į	II. N	on-establishment of opinion with regard to novelty, inventive step and industrial applicability			
	1. T	The questions whether the claimed invention appears to be novel, to involve an inventive step (to be non-obvious), or to be industrially applicable have not been examined in respect of:			
	☐ the entire international application,				
	Ø	claims Nos. 10-12,18			
		because:			
	⋈	the said international application, or the said claims Nos. 18 relate to the following subject matter which does not require an international preliminary examination (specify):			
		see separate sheet			
	Ø	the description, claims or drawings (indicate particular elements below) or said claims Nos. 10-12 are so unclear that no meaningful opinion could be formed (specify):			
		see separate sheet			
		the claims, or said claims Nos. are so inadequately supported by the description that no meaningful opinion could be formed.			
		no international search report has been established for the said claims Nos.			
2		neaningful international preliminary examination cannot be carried out due to the failure of the nucleotide and/ amino acid sequence listing to comply with the standard provided for in Annex C of the Administrative tructions:			
	☐ the written form has not been furnished or does not comply with the Standard.				
		the computer readable form has not been furnished or does not comply with the Standard.			
IV	. La	ck of unity of invention			
1.	ln r	esponse to the invitation to restrict or pay additional fees, the applicant has:			
		restricted the claims.			
	×	paid additional fees.			
		paid additional fees under protest.			
		neither restricted nor paid additional fees.			
2.		This Authority found that the requirement of unity of invention is not complied with and chose, according to Rule 68.1, not to invite the applicant to restrict or pay additional fees.			
3.	This	Authority considers that the requirement of unity of invention in accordance with Rules 13.1, 13.2 and 13.3			
		complied with.			
		not complied with for the following reasons:			

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4.	Consequently, the following parts of the international application were the subject of international preliminary examination in establishing this report:
	- •

☐ all parts.

oxtimes the parts relating to claims Nos. 1-9,13-17,19-22 (all part) .

V. Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

1. Statement

Novelty (N)

Yes: Claims

1-9,13-17,19-22 (all part)

No: Claims

Inventive step (IS)

Yes: Claims

No: Claims

1-9,13-17,19-22 (all part)

Industrial applicability (IA)

Yes: Claims

1-9,13-17,19-22 (all part)

No: Claims

2. Citations and explanations

see separate sheet

SECTION III

- Claim 18 relates to the treatment of human and/or animal bodies. According to 1). Rule 67(1)(iv) an examination is not required for such a claim.
- 2). The Applicant is asked to explain the reason for the provisos found in claim 1. If they are intended to exclude some unacknowledged prior art known to the Applicant the said prior art should be cited and made available.

SECTION IV

- 1). Relevant prior art is represented by:
- D1 WO-A-02/26722
- 2). An international patent application can contain only one invention or a plurality of inventions, if they are linked together by a single inventive concept.

In other word, there must be a specific technical feature common to all the claimed alternatives and which makes a contribution over the prior art taken as a whole.

Although all the claimed compounds seem to have a common pharmaceutical activity (antiinflammatory), a specific technical element shared by all the alternatives and making this contribution cannot be identified.

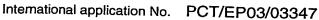
D1 discloses also compounds having the same pharmaceutical properties as those of the current application and (see example 60) discloses also compounds falling within the claimed scope.

It has to be noted that the mere disclaiming of such a compound cannot restore unity of invention.

The separate inventions/groups of invention are:

- Claims 1-22 in which R¹ is imidazolyl 1.
- 2. Claims 1-22 in which R1 is triazolyl

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- 3. Claims 1-22 in which R1 is oxadiazolyl
- 4. Claims 1-22 in which R1 is thiazolyl
- 5. Claims 1-22 in which R1 is thiophenyl
- 6. Claims 1-22 in which R1 is isoxadiazolyl
- 7. Claims 1-22 in which R1 is isoxathiazolvl
- 8. Claims 1-22 in which R1 is pyridinyl
- 9. Claims 1-22 in which R1 is furanyl
- Claims 1-22 in which R1 is isoxazolyl 10.
- 11. Claims 1-22 in which R1 is tetrazolyl
- Claims 1-22 in which R1 is pyrazolyl 12.

The applicant paid the corresponding fees for the inventions in which R1 is furanyl, oxadiazolyl and pyrazolyl. This opinion is therefore limited to these subject-matters.

SECTION V

- 1). Relevant prior art is represented by:
- D1 WO-A-02/26722
- **D2** EP-A-243959
- D3 J.Med.Chem. (1991), vol. 34, p. 616-24
- D4 WO-A-00/71518
- The claimed matter is novel vis-à-vis D2 and D3, since the grouping "Y" is a single 2). bond for the compounds disclosed in these documents.

The claimed matter is a selection vis-à-vis D4 but it is regarded as novel, since the specific combination of the structural elements of the claimed compounds is not disclosed in D4 (values of Y and values of R1).

None of the cited documents discloses compounds having antiinflammatory 3). properties as those currently claimed.

Thus, the problem underlying the current application appears to be the provision of further morpholinyl derivatives having antiinflammatory properties.

The data of the description show that this problem has been solved by some of the claimed

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compounds.

In the absence of any relevant prior art, the skilled person would not arrive at the claimed compounds by using only his technical knowledge.

An inventive step is therefore not acknowledged on the whole claimed scope, since the wording of the claims contains expressions which are unlimited and therefore lead to an unlimited number of compounds which inherently cannot represent a solution to the given problem.

4). There is no objection with regard to industrial applicability.